

K123805 E-Z CHEK BLOOD LEAK TEST STRIPSMar 20, 2013
99 days to decisionK123805 · Product code: **FJD** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k123805/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector, Leak, Blood (FJD)
Date received	Dec 11, 2012
Decision date	Mar 20, 2013
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Reprocessing Products Corp
Location	Tucson, AZ, US
Contact	TED WILLIAMS
510(k) history	6 submissions · 6 cleared · 1995-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k123805/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 9, 2026